#### **PCT**

(30) Priority Data: 07/997,281

# WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



#### INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

| (51) International Patent Classification 5:           |          | (11) International Publication Number:  | WO 94/15219            |  |
|---|----------|---|------------------------|--|
| G01N 35/00  | A1       | (43) International Publication Date:  | 7 July 1994 (07.07.94) |  |
| (21) International Application Number: PCT/US93/12424 |          | (81) Designated States: AT, AU, BB, BG, BR, BY, CA, CH, CDE, DE, DK, ES, FI, GB, HU, JP, KP, KR, KZ, LK, LU, L' |                        |  |
| (22) International Filing Date: 20 December 1993      | (20.12.9 | 3) MG, MN, MW, NL, NO, NZ, PL   | PT. RO, RU, SD, SE,    |  |

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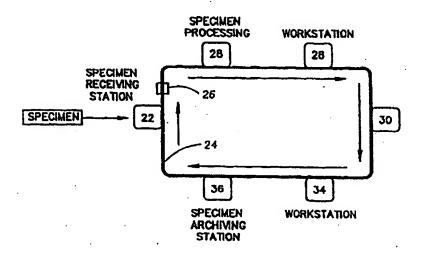
(74) Agents: FREDERIKSEN, Mark et al.; Zarley, McKee, Thomte, Voorhees & Sease, 801 Grand Avenue, Suite 3200, Des Moines, IA 50309 (US). 51) Designated States: AT, AU, BB, BG, BR, BY, CA, CH, CZ, DE, DK, ES, FI, GB, HU, IP, KP, KR, KZ, LK, LU, LV, MG, MN, MW, NL, NO, NZ, FL, FT, RO, RU, SD, SE, SK, UA, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BP, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).

Published

With international search report.

(54) Title: METHOD FOR AUTOMATIC TESTING OF LABORATORY SPECIMENS

23 December 1992 (23.12.92)



(57) Abstract

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A method for automatic testing of a laboratory specimen includes the initial step of obtaining a specimen to be tested and placing the specimen in a specimen container. The container is removably mounted in an independent carrier (26) designed to carry an individual specimen through a laboratory to one or more of a plurality of work stations (28, 30, 32, 34), where a predetermined test will be performed on the specimen. Once the test has been performed, the carrier (26) is moved to an archiving station (36) for storage of the specimen.

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#### METHOD FOR AUTOMATIC TESTING OF LABORATORY SPECIMENS

#### Technical Field

The present invention relates generally to laboratory automation systems, and more particularly to an improved method for automating a laboratory for the testing of individual laboratory specimens.

#### Background of the Invention

Clinical laboratory testing has changed and improved remarkably over the past 70 Initially, tests or assays were performed manually, and generally utilized large quantities of serum, blood or other materials/body fluids. As mechanical technology developed in the industrial work place, similar technology was introduced into the clinical With laboratory. the introduction technology, methodologies were also improved in an effort to improve the quality of the results produced by the individual instruments, minimize the amount of specimen required to perform each test.

More recently, instruments have been developed to increase the efficiency of testing procedures by reducing turn around time and decreasing the volumes necessary to perform various assays. Present directions in laboratory testing focus on cost containment procedures and instrumentation. Laboratory automation is one area in which cost containment procedures are currently being explored. Robotic engineering has evolved to such a degree that various types of robots have been applied in the clinical laboratory setting.

focus main of prior art automation relies on the implementation of conveyor systems to connect areas of the clinical laboratory. Known conveyor systems in the laboratory setting utilize separate conveyor segments to move specimens from a processing station to a specific laboratory work station. In order to obtain cost savings, the specimens are sorted manually, and grouped in a carrier rack to be conveyed to a specific location. In this way, a carrier will move a group of 5-20 specimens from the processing location to the specific work station to perform a single test on each of the specimens in the carrier.

While grouping a plurality of specimens in a single carrier may be more cost efficient where every specimen requires only a single specific test, and none of the specimens within a carrier require special priority, it is not uncommon in the hospital environment for a specimen to be subjected to a variety of different tests, or for a particular specimen to require a very short turn around time. In such an event, the current automation system could not be utilized, and the particular specimen

would have to be manually moved to various work test stations based upon the time constraints and tests designated for the specimen.

Another problem with prior attempts at laboratory automation is in tracking the specimen and reporting the results of the specimen tested. Test results can serve as the basis for requiring additional testing of a particular specimen reflex or spawned testing. If the test results are required within a short time period, rapid and efficient reporting of test results can improve laboratory quality and efficiency.

#### Summary of the Invention

It is a general object of the present invention to provide a method for automating a clinical laboratory which permits individual and independent assignment of a specimen to one or more of a plurality of work stations within the laboratory.

Another object of the present invention is to provide a method for automating a clinical laboratory which can improve turn around time for the testing of an individual specimen.

Still another object is to provide a method for automating a laboratory which permits automatic conveyance of a specimen to a plurality of work stations.

Still another object of the present invention is to provide a method for automating a clinical laboratory which tracks a specimen location throughout the laboratory and reports test results to a central database for immediate review by a doctor. These and other objects will be apparent to those skilled in the art.

The method for automatic testing of a laboratory specimen of the present invention includes the initial step of obtaining a specimen to be tested and placing the specimen in a specimen container. The container is removably mounted in an independent carrier designed to carry an individual specimen of a number of different sizes and shapes through a laboratory to one or more of a plurality of work where a predetermined test will stations, performed on the specimen. Once the carrier has arrived at the predetermined work station, the carrier is removed from the conveyor and a test is The carrier is then conducted on the specimen. returned to the conveyor and moved to an archiving Preferably, a station for storage of the specimen. computer is incorporated with the laboratory work stations, and includes a sensor located at each work station and archiving station. Each carrier and specimen container is marked with an identification code which is read by the sensor and transmitted to The computer may then operate a the computer. carrier removal apparatus at a predetermined work station to remove the carrier at the appropriate location for testing. Keyboards located at each work station permit the entry of test results at the After a particular test has been work stations. completed, the carrier is placed on the conveyor once again, and may be directed to an additional work station or to the archiving station. conveyor system is preferably arranged in a closed loop formation such that a specimen can be moved to any specific work station in any specific sequence.

#### Brief Description of the Drawings

Figure 1 is a flow chart showing the integration of a laboratory automation system with a laboratory information system and hospital information system;

Figure 2 is a schematic diagram of specimen movement through a laboratory automation system; and

Figure 3 is an enlarged schematic view of the specimen processing station and one work station along the schematic of Figure 2.

#### Description of the Preferred Embodiment

Referring now to the drawings, Figure 1 is a flowchart showing how the laboratory automation system (LAS) of the present invention integrates with the day-to-day operations of a hospital. Box 10 refers to any patient who is in need of examination and/or diagnosis. Box 12 represents the relevant physician or other practitioner who will interpret the results

of the examination in order to determine the necessity of tests, in order to make a final diagnosis and/or prescribe a specified treatment. Information passes in both directions between doctor and patient during this examination.

As a result of the examination, the doctor will make a record of the examination results, and may enter a request for a specific test to be performed. This information is entered in the general hospital information system (HIS) shown as box 14 in the flowchart. The HIS will correlate patient identification information, room information, as well as any insurance or other typical general information necessary for operation of a hospital.

The HIS is a computer system which communicates with various areas of the hospital to integrate all functions of the hospital.

Once the doctor's test order is correlated with the patient identification information, the HIS will forward the correlated information to the laboratory information system (LIS) designated as box 16 in the Figure 1. The LIS is a computer system which is connected to the HIS to quickly and efficiently communicate information.

As shown in Figure 1, the LIS assigns the task specimen to an appropriate obtaining а of technician, the retrieval of the specimen designated generally at box 18. The physical specimen obtained from the patient is then entered in the laboratory automation system (LAS) designated generally as box The LAS takes the place of prior art manual testing procedures, including the reporting of the test results to the LIS. The LIS communicates with the LAS to order specific tests related to a specific specimen, and receive the results of those The LIS also communicates with the HIS to report test results for accounting and insurance The LIS reports either to the doctor via a separate work station, or via the HIS, to report the results of the requested tests.

Referring now to Figure 2, a schematic diagram of specimen movement throughout the laboratory automation system is shown. The specimen arrives at a specimen receiving station 22, where the specimen is entered on a conveyor system designated generally at 24. During the assignment of the task of obtaining a specimen, the laboratory information system would also provide a specimen container

marked with an appropriate patient identification The inventor has found that a conventional bar code label applied to the specimen container is a simple and efficient method for fulfilling this function. Since most specimen containers are not designed for transport on a conveyor system, separate carrier 26 is provided to support an individual specimen container on conveyor system 24. At specimen receiving station 22, the carrier 26 is given an identification code which correlates with the specimen container, so that the container and carriage may be directed throughout the laboratory automation system, even when the specimen container is removed from the carriage for specific testing at a work station.

As shown in Figure 2, conveyor system 24 is preferably a continuously moving conveyor which will move carriers 26 in a generally closed loop system. The first station which a carrier 26 will encounter after entry on conveyor system 24, is specimen processing station 28. At processing station 28, the carrier assignment is entered into the LAS to determine which work stations the specimen must utilize, the order in which the stations are to be utilized and any other pertinent information with respect to priority or turn around time.

While Figure 2 shows only 3 specific work stations, 30, 32, and 34, obviously a conventional clinical laboratory could have a wide variety of such stations throughout a facility. The closed loop system of conveyor 24 permits a specimen to stop at any given work station in any particular order. Thus, if time constraints require that the test of work station 34 be performed first, and that

a test of work station 32 be performed at some time after the test of work station 34, the specimen can travel on conveyor 24 past work stations 30 and 32, directly to work station 34, for immediate testing. Carrier 26 is then reintroduced on conveyor system 24 to follow the closed loop around to the next work station assigned to the specimen. Once the testing has been completed, the specimens are forwarded to the specimen archiving station 36 for removal from conveyor 24 and appropriate storage.

Referring now to Figure 3, an enlarged view of a portion of the schematic of Figure 2 is shown. Specimen processing station 28 and work station 30 are shown in schematic view to demonstrate each specific work station located along conveyor system As carrier 26 moves along conveyor 24, it will pass within the zone of specimen processing station 28 where a sensor 38 will detect the identification code on carrier 26. In the preferred embodiment of the invention, sensor 38 is a bar code reader while the identification code on the carrier 26 is a bar Sensor 38 is connected with the LIS, to In the example record the movement of carrier 26. of Figure 3, carrier 26 has just entered the conveyor system 24, and therefore will be assigned to stop at the specimen processing station 28.

A gate 40 is connected to the LIS and will be activated to redirect the movement of carrier 26 off of conveyor 24 and on to an auxiliary conveyor 42 to reach the ultimate processing location 28a within the processing station 28. Processing area 28a may be comprised of manual processing, or fully automatic mechanical processing. An additional sensor 44 is positioned along auxiliary conveyor 42

to track the location of the carrier and specimen, and may be utilized to activate any automatic mechanical equipment associated with the specimen processing work area 28a.

discussed above, the specimen processing station is utilized to direct the movement of the specimen to the appropriate work station at the A keyboard 29, or the like, is appropriate time. provided to enter the information into the LIS. This information is downloaded to the LIS which in turn distributes the appropriate instructions to the pertinent sensors and work stations, as described in more detail hereinbelow. Once processing has been completed, the specimen is again loaded in specimen carrier 26 and placed in conveyor system 24 by auxiliary conveyor 42. This procedure can be accomplished by virtue of sensor 44 or manually within the work area 28a of processing station 28.

In the present example, work station 30 has been designated as the first testing area for the specimen. Thus, conveyor 24 will move specimen carrier 26 into the zone of work station 30. A sensor 38' will acknowledge the passage of carrier 26 thereby, thereby triggering the LIS to direct gate 40' to divert the carrier 26 onto the auxiliary conveyor 42' of work station 30. A sensor 34' will then direct the specimen to the appropriate testing area 30a.

Once the test performed by work station 30 has been completed, the results are transmitted from the work area 30a to the LIS by virtue of keyboard 31, and the specimen is loaded in the specimen carrier 26 and positioned on auxiliary conveyor 42'. The specimen will then be moved to the main conveyor

system 24 for movement to the next appropriate station. Work stations 32 and 34 are not shown in detail, but include the same basic equipment as work station 30. Thus, a sensor 38' located at work stations 32 and 34 will acknowledge passage of the specimen at that location and either direct the specimen into the work station, or direct the specimen to continue past the work station. order in which the tests are conducted is important, the specimen can be directed to bypass any work station along the conveyor system 24 so as to immediately reach the highest priority work station to perform the appropriate testing. Since the conveyor system is a closed loop, the specimen can then be moved around the loop to any other work station.

Once all requested tests have been performed, the specimen will be directed into the specimen archiving station utilizing a sensor 38' and gate 40' in the same manner as work stations 30, 32 and Since every sensor 38, 38', 44 and 44' interconnected by way of the laboratory information system, the location and status of any specimen is always readily accessible by the doctor. Since the the doctor can call is programmable, additional tests at any time during the movement of the specimen within the LAS. This ability to direct an individual specimen to one or more of a plurality of work stations decreases the turn around time and increases the versatility of the automation system. With the use of robotics, and a fully integrated laboratory instrumentation, it is possible to fully automate the entire laboratory automation system. In addition, the results of standard testing may conventionally require additional testing. In such a case, the LIS may automatically assign additional or different work station stops based upon the results received from a test at any given work station. The capability of prioritizing the testing, also permits a doctor to diagnose and/or otherwise individualize the test battery which is required for an individual patient.

Whereas the invention been has shown described in connection with the preferred embodiment thereof, it will be understood that many modifications, substitutions and additions may be made which are within the intended broad scope of the appended claims. There has therefore been shown and described an improved method for testing of laboratory specimen.

I claim:

1.

A method for automatically testing a specimen in a laboratory, comprising the steps of:

- providing a specimen to be tested and placing the specimen in a specimen container;
- providing a laboratory with a plurality of work stations, each work station adapted to conduct a predetermined test on a specimen, and an operable conveyor extending between said work stations;
- providing independent carriers for carrying an individual specimen container along said conveyor;
- placing said specimen container in a carrier and placing the carrier on the conveyor;
- operating said conveyor to move the carrier to a predetermined one of said plurality of work stations;
- removing the carrier from the conveyor at said predetermined work station;
- conducting a predetermined test on said specimen at said work station;
- returning the carrier to said conveyor upor completion of the test; and
- operating said conveyor to move said first carrier to an archiving station for storage of specimens.

2.

The method of claim 1, further comprising the step of marking each of the specimen container and carrier with an identification code, prior to the step of placing the carrier on the conveyor.

3.

The method of claim 2, further comprising the step of providing a receiving station along said conveyor, and wherein the marking step is performed at said receiving station.

4.

The method of claim 3, further comprising the step of providing a sensor means at each work station and said receiving station for reading said identification code on said carrier as the carrier moves on the conveyor at the work stations and receiving station.

5.

The method of claim 4, further comprising the step of providing a computer connected to said sensors and having a central database with stored information regarding said specimen, and including the step of said sensors transmitting the identification code read from the carrier to said database as the carrier moves past each said sensor.

6.

The method of claim 5, further comprising the steps of:

apparatus at each work station providing removing the carrier from the conveyor, each said removing apparatus being connected to said computer and operable in response to commands from said computer, said computer programmed to the removing apparatus command predetermined work station to remove the carrier from the conveyor in response to a signal the sensor at said predetermined work station;

said sensor at said predetermined work station transmitting a signal to the computer identifying the carrier as the carrier moves past the sensor; and

said computer commanding the removal apparatus at the predetermined work station to remove the carrier from the conveyor, in response to said sensor signal.

7,

The method of claim 6, further comprising the step of providing computer input means at each said work station, connected to said computer, and the step of entering results of said predetermined test into the computer upon completion of said test.

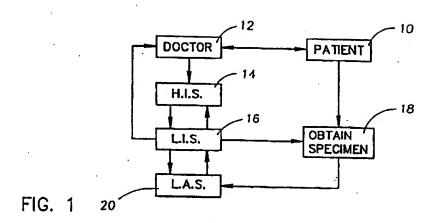
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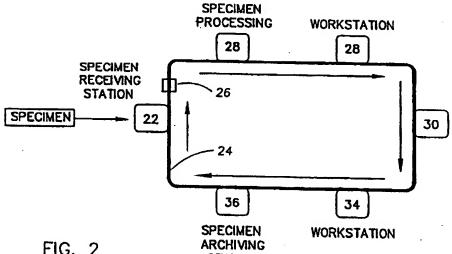
The method of claim 7, further comprising the steps of:

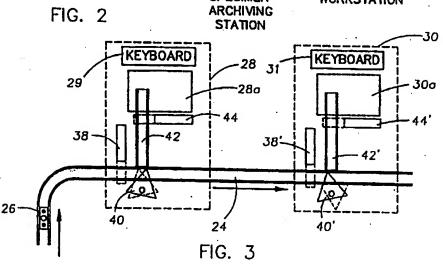
- said computer being programmed to command the removing apparatus at said archiving station to remove the carrier from the conveyor upon the occurrence of two events: (a) the step of
- returning the carrier to the conveyor upon completion of the test, and (b) receipt of a signal from the archiving station sensor;
- said archiving station sensor transmitting a signal to the computer identifying the carrier as the carrier moves past the sensor; and
- said computer commanding the archiving station removal apparatus to remove the carrier from the conveyor, in response to said sensor signal and after the step of returning the carrier to the conveyor after completion of the test.

9.

The method of claim 1, wherein said conveyor is a continuously moving conveyor arranged in a closed loop such that a carrier will continuously move along the conveyor until removed therefrom.







#### INTERNATIONAL SEARCH REPORT

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|   |  |   |                         |
| Υ   | CHEMOMETRICS & INTELLIGENT LABOR   | ATORY   | 1-9                     |
|   | SYSTEMS,   |   |                         |
| ·   | vol.17, no.1, October 1992, AMST   | FERDAM NL   |                         |
|   | pages 111 - 118, XP00032° `3<br>& FIRST INTERNATIONAL S' JIUM              | ON  |                         |
|   | AUTOMATION, ROBOTICS /. ATIFIC   |   |                         |
|   | INTELLIGENCE APPLIED TO ANALYTIC   | AL  |                         |
|   | CHEMISTRY, MONTREUX, SWITZERLAND   | ), 26-28  |                         |
|   | FEB. 1992,, ISSN 0925-5281   | nebettes  |                         |
| l   | Felder R A 'Clinical laboratory in the 1990s'                              | ropotics  |                         |
| ]   | see page 113, right column, para   | igraph 2 -  |                         |
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| j   | see page 115, right column, para   | agraph 2  |                         |
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| * Special c   | stegaries of cited documents:  | "T" later document published after the m  | ternational filing date |
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| considered to be of particular relevance invention  "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention  |  |   | c claimed invention     |
| filing date  Cannot be considered novel or cannot be considered from the document is taken alone  "I." document which may throw doubts on priority claim(s) or involve an inventive step when the document is taken alone           |  |   | ot be considered to     |
| which is clied to establish the publication date of another catation or other special reason (as specified)  "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the |  |   | e darmed invention      |
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| later   | than the priority date claimed   | '&' document member of the same pater   |                         |
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| ;   | 7 April 1994   | 1 3. 04. 9  | 4                       |
| Name and  | mailing address of the ISA   | Authorized officer  |                         |
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#### INTERNATIONAL SPARCH REPORT

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|           | tion) DOCUMENTS CONSIDERED TO BE RELEVANT  | I Delevere se elejes No |
|-----------|--|-------------------------|
| ategory * | Citation of document, with indication, where appropriate, of the relevant passages                         | Relevant to claim No.   |
| ,         | DE,A,39 34 890 (OLYMPUS OPTICAL CO) 26<br>April 1990   | 1-9                     |
|           | see column 4, line 12 - line 24<br>see column 6, line 54 - column 10, line 9;                              |                         |
|           | figures 3-5  |                         |
|           | EP,A,O 417 006 (TOA MEDICAL ELECTRONICS) 13 March 1991   | 1                       |
|           | see column 20, line 41 - column 22, line<br>14; figure 5   | ·                       |
|           | HITACHI REV. (JAPAN), HITACHI REVIEW,<br>TOKYO, JP,  | 1-7                     |
|           | vol.41, no.4, September 1992, ISSN 0018-277X   |                         |
|           | pages 167 - 172, XP000334656  Ikeda T et al 'Total clinical laboratory                                     |                         |
|           | testing system for laboratory automation' see page 169 - page 170; figures 4-6                             |                         |
|           | FR.A.2 600 166 (RHONE POULENC RECHERCHES) 18 December 1987   | ·                       |
| ι.        | DE,B,25 19 111 (SIEMENS AG) 16 July 1976   |                         |
| γ,χ       | CHEMOMETRICS & INTELLIGENT LABORATORY SYSTEMS, vol.21, no.2/3, 8 December 1993,                            | 1-9                     |
|           | AMSTERDAM NL pages 169 - 179, XP000413289  |                         |
|           | & SECOND INTERNATIONAL SYMPOSIUM ON AUTOMATION, ROBOTICS AND ARTIFICIAL INTELLIGENCE APPLIED TO ANALYTICAL |                         |
|           | CHEMISTRY, MONTREUX, SWITZERLAND, 23-26 FFB. 1993. ISSN 0925-5281  | ·                       |
|           | R.S. MARKIN 'Implementing automation in a modern clinical laboratory'                                      |                         |
|           | see the whole document   | 1-9                     |
| P,Y       | US,A,5 224 585 (E.E. BLANCO ET AL.) 6 July 1993  |                         |
| ٠.        | see column 3, line 28 - line 68; figures 1,3-8   |                         |
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#### INTERNATIONAL SEARCH REPORT

Information on patent family members

Intern si Application No
PCT/US 93/12424

| Patent document cited in search report | Publication date | Patent family member(s)                            |   | Publication date   |
|--|------------------|--|---|--|
| DE-A-3934890                           | 26-04-90         | JP-A-<br>JP-A-<br>US-A-                            | 2110376<br>2163660<br>5087423                                 | 23-04-90<br>22-06-90<br>11-02-92                                     |
| EP-A-0417006                           | 13-03-91         | JP-A-<br>AU-B-<br>AU-A-<br>US-A-<br>US-A-<br>AU-A- | 3094159<br>636384<br>6213090<br>5270012<br>5209903<br>3198393 | 18-04-91<br>29-04-93<br>14-03-91<br>14-12-93<br>11-05-93<br>01-04-93 |
| FR-A-2600166                           | 18-12-87         | US-A-  | 4805469   | 21-02-89   |
| DE-B-2519111                           | 15-07-76         | .NONE  |   |  |
| US-A-5224585                           | 06-07-93         | AU-8-<br>WO-A-                                     | 3813893<br>9320940  | 18-11-93<br>28-10-93   |

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